

GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: October 6, 2002, 12:40:30 : Search time 284 Seconds
(without alignments)
120.909 Million cell updates/sec

Title: US-09-754-468-47
Perfect score: 20
Sequence: 1 gattagcataaataatc 20

Scoring table: IDENTITY_NUC
Gapop 10.0, Gapex 1.0

Searched: 1736436 segs, 858457221 residues
Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Geneseq_032802.1
1: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA1980.DAT:*
2: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA1981.DAT:*
3: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA1982.DAT:*
4: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA1983.DAT:*
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9: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA1988.DAT:*
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13: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA1992.DAT:*
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15: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA1994.DAT:*
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19: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA1998.DAT:*
20: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA1999.DAT:*
21: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA2000.DAT:*
22: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA2001A.DAT:*
23: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA2001B.DAT:*
24: /net/abs06/SIDSL1/gcgdata/hold-geneseq/geneseqn-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20	100.0	836	AAH25706	E coli secA coding
2	20	100.0	836	AAH25707	E coli secA mRNA a
3	20	100.0	2754	AAH38420	JPL1192089 Seg ID
4	20	100.0	3811	AAH38421	E. coli secA DNA.
5	18.4	92.0	325	AAH05228	Human secreted pro
6	18.4	92.0	645	AAH61816	Human immune/haema
7	18.4	92.0	1972	AAH81074	Human immune/haema
8	18.4	92.0	1972	AAH81075	Human immune/haema
9	18.4	92.0	1976	AAH81076	Human immune/haema

C 10	17.4	87.0	3153	21	AAH59841	Human secreted pro
C 11	16.4	82.0	401	22	AAH95332	Human neutregulin g
C 12	16.4	82.0	401	22	AAH95333	Human neutregulin g
C 13	16.4	82.0	401	22	AAH95334	Human neutregulin g
C 14	16.4	82.0	401	22	AAH95335	Human neutregulin g
C 15	16.4	82.0	401	22	AAH95336	Human neutregulin g
C 16	16.4	82.0	401	22	AAH95337	Human neutregulin g
C 17	16.4	82.0	401	22	AAH95338	Human neutregulin g
C 18	16.4	82.0	401	22	AAH95339	Human neutregulin g
C 19	16.4	82.0	401	22	AAH95340	Human neutregulin g
C 20	16.4	82.0	401	22	AAH95341	Human neutregulin g
C 21	16.4	82.0	401	22	AAH95342	Human neutregulin g
C 22	16.4	82.0	401	22	AAH95343	Human neutregulin g
C 23	16.4	82.0	401	22	AAH95344	Human neutregulin g
C 24	16.4	82.0	401	22	AAH95345	Human neutregulin g
C 25	16.4	82.0	401	22	AAH95346	Human neutregulin g
C 26	16.4	82.0	401	22	AAH95347	Human neutregulin g
C 27	16.4	82.0	401	22	AAH95348	Human neutregulin g
C 28	16.4	82.0	401	22	AAH95349	Human neutregulin g
C 29	16.4	82.0	401	22	AAH95350	Human neutregulin g
C 30	16.4	82.0	401	22	AAH95351	Human neutregulin g
C 31	16.4	82.0	401	22	AAH95352	Human neutregulin g
C 32	16.4	82.0	401	22	AAH95353	Human neutregulin g
C 33	16.4	82.0	401	22	AAH95354	Human neutregulin g
C 34	16.4	82.0	401	22	AAH95355	Human neutregulin g
C 35	16.4	82.0	401	22	AAH95356	Human neutregulin g
C 36	16.4	82.0	401	22	AAH95357	Human neutregulin g
C 37	16.4	82.0	401	22	AAH95358	Human neutregulin g
C 38	16.4	82.0	401	22	AAH95359	Human neutregulin g
C 39	16.4	82.0	401	22	AAH95360	Human neutregulin g
C 40	16.4	82.0	401	22	AAH95361	Human neutregulin g
C 41	16.4	82.0	401	22	AAH95362	Human neutregulin g
C 42	16.4	82.0	401	22	AAH95363	Human neutregulin g
C 43	16.4	82.0	401	22	AAH95364	Human neutregulin g
C 44	16.4	82.0	401	22	AAH95365	Human neutregulin g
C 45	16.4	82.0	401	22	AAH95366	Human neutregulin g

ALIGNMENTS

AAH25706/C	standard; DNA: 836 BP.
AAH25706;	
14-AUG-2001 (first entry)	
E coli secA coding sequence.	
Antisense; microbial growth; essential gene; antimicrobial;	
proliferation; infectious disease; secA; ds.	
Escherichia coli.	
US6228579-B1.	
08-MAY-2001.	
14-NOV-1997;	9705-0971090.
14-NOV-1997;	9705-0971090.
(UYSA-) UNIV SAN DIEGO STATE FOUND.	
Zyskind JW, Forsyth RA;	
WPI, 2001-335011/35.	
Identifying microbial proliferation genes; useful for identifying	
antimicrobial agents, comprises introducing into a microorganism an	
exogenous nucleic acid having sequence identity to an endogenous	

PT microbial gene -

PS Disclosure: Fig 11: 28pp; English.

CC The present invention describes a method of identifying genes essential
CC for microbial growth and proliferation, involving introducing an
CC exogenous nucleic acid into a microorganism, where the sequence is
CC similar to an endogenous microbial gene, and identifying the gene as
CC essential by comparing the organism's viability when the exogenous
CC sequence is expressed and when it is not present. This can be used to
CC identify targets for antimicrobial compounds for use in the therapy of
CC infectious diseases.

SO Sequence 836 BP; 223 A; 220 C; 198 G; 195 T; 0 other;

Query Match

Best Local Similarity 100.0%; Score 20; DB 22; Length 836;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTAGCATATTAATAATCTC 20

DB 648 GATTAGCATATTAATAATCTC 629

RESULT 2

AAH25707

ID AAH25707 standard; RNA; 836 BP.

AC AAH25707;

DT 14-AUG-2001 (first entry)

DE E coli secA mRNA antisense sequence.

KW Antisense: microbial growth; essential gene; antimicrobial;

KM proliferation; infectious disease; secA; ss.

OS Escherichia coli.

XX US6228579-B1.

XX 08-MAY-2001.

XX 14-NOV-1997; 97US-0971090.

XX 14-NOV-1997; 97US-0971090.

PA (URSA-) UNITV SAN DIEGO STATE FOUND.

XX Zyskind JW, Forsyth RA.

XX WPI: 2001-335011/35.

PT Identifying microbial proliferation genes, useful for identifying
PT antimicrobial agents, comprises introducing into a microorganism an
PT exogenous nucleic acid having sequence identity to an endogenous
PT microbial gene -

Example 3: Fig 12; 28pp; English.

CC The present invention describes a method of identifying genes essential
CC for microbial growth and proliferation, involving introducing an
CC exogenous nucleic acid into a microorganism, where the sequence is
CC similar to an endogenous microbial gene, and identifying the gene as
CC essential by comparing the organism's viability when the exogenous
CC sequence is expressed and when it is not present. This can be used to
CC identify targets for antimicrobial compounds for use in the therapy of
CC infectious diseases.

SO Sequence 836 BP; 195 A; 198 C; 220 G; 223 U; 0 other;

Query Match

Best Local Similarity 100.0%; Score 20; DB 22; Length 836;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Matches 14; Conservative 6; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTAGCATATTAATAATCTC 20

DB 189 GAUAGCAUAAUAAUCC 208

RESULT 3

AAH8420/c

ID AAH8420 standard; DNA; 2754 BP.

AC AAH8420;

DT 30-SEP-1999 (first entry)

DE JP1192089 Seq ID 4.

KW Tp surface antigen; secretion-related enzyme; ds.

OS Unidentified.

XX JP1192089-A.

XX 21-JUL-1999.

XX 29-DEC-1997; 97JP-0367638.

XX 29-DEC-1997; 97JP-0367638.

PA (PURE) FUJIREBIO KK.

XX WPI: 1999-461459/39.

PT Treponema pallidum-fused DNA sequence - and expression of Treponema
PT pallidum antigen by using said sequence

PS Disclosure: Page 10-12; 16pp; Japanese.

CC This invention describes a novel Treponema pallidum (Tp)-fused DNA
CC sequence in which a DNA sequence coding the surface antigen of Tp added
CC by a signal peptide is fused with a DNA sequence coding a
CC secretion-related enzyme. Also claimed is a method for expressing the
CC Tp antigen by using the above Tp-fused DNA sequence.

SO Sequence 2754 BP; 729 A; 680 C; 761 G; 584 T; 0 other;

Query Match

Best Local Similarity 100.0%; Score 20; DB 20; Length 2754;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTAGCATATTAATAATCTC 20

DB 49 GATTAGCATATTAATAATCTC 30

RESULT 4

AAH38291/c

ID AAH38291 standard; DNA; 3811 BP.

AC AAH38291;

DT 16-JUN-1999 (first entry)

DE E. coli secA DNA.

KW Microorganism inhibitor; antisense; nuclease resistant; treatment;

KM ribonucleotide reductase; secA gene; pathological condition;

KW antimicrobial agent; crop protection; ss.

OS Escherichia coli.

XX WO9902673-A2

PD 21-JAN-1999.
XX 10-JUL-1998: 98WO-CA00666.
PF 10-JUL-1997: 97US-0052160.
XX (GENE-) GENESENSE TECHNOLOGIES INC.
XX Dugourd D, Wright JA, Young AH;
XX WPI; 1999-120874/10.
DR
XX
XX New oligonucleotides complementary to RR or SecA genes - useful to
XX inhibit growth of microorganisms
XX
XX Disclosure; Fig 5; 103pp; English.
XX
XX This invention describes novel antisense oligonucleotides
CC (AA38301-X38552) which are nuclease resistant, and comprises about 3-50
CC nucleotides complementary to the ribonucleotide reductase gene or the
CC secA gene of a microorganism. The antisense oligonucleotides are used to
CC treat mammalian pathological conditions mediated by microorganisms. The
CC oligonucleotides are particularly useful as antimicrobial agents in crop
CC protection. This DNA sequence contains the Escherichia coli secA gene.
XX
SQ Sequence 3811 BP; 1016 A; 942 C; 1000 G; 853 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 3811;
Best Local Similarity 100.0%; Pred. No. 7.3;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTAGCATATATAAATCTC 20
DB 830 GATTAGCATATATAAATCTC 811

RESULT 5
AAC05228/C
ID AAC05228 standard; cDNA; 325 BP.
XX
XX AAC05228;
XX
XX 06-OCT-2000 (first entry)
XX
DE Human secreted protein 5' EST, SEQ ID NO: 9303.
XX
XX Human: 5' EST; expressed sequence tag; secreted protein; cDNA isolation;
XX gene therapy; chromosome mapping; ss.
XX
XX Homo sapiens.
XX
XX EP1033401-A2.
XX
XX 06-SEP-2000.
XX
XX 21-FEB-2000; 2000EP-0200610.
XX
XX 26-FEB-1999; 99US-0122487.
XX
XX (GEST) GENSET.
XX
XX Dumas Milne Edwards J, Duclert A, Giordano J;
XX WPI; 2000-500381/45.
XX
XX New nucleic acid that is a 5' expressed sequence tag (5' EST) for
XX obtaining cDNAs and genomic DNAs that correspond to 5' ESTs and for
XX diagnostic, forensic, gene therapy and chromosome mapping procedures -
XX
XX Claim 1: SEQ ID 9303; 71pp + CD-ROM; English.
XX
XX The present sequence is one of a large number of 5' ESTs derived from
CC mRNAs encoding secreted proteins. No ORF has yet been conclusively

CC identified within the present sequence. The 5' ESTs were prepared from
CC total human RNAs or polyA+ RNAs derived from 30 different tissues. EST
CC sequences usually correspond mainly to the 3' untranslated region (UTR)
CC of the mRNA because they are often obtained from oligo-dT primed cDNA
CC libraries. Such ESTs are not well suited for isolating cDNA sequences
CC derived from the 5' ends of mRNAs and even in those cases where longer
CC cDNA sequences have been obtained, the full 5' UTR is rarely included.
CC 5' ESTs are derived from mRNAs with intact 5' ends and can therefore be
CC used to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used
CC in diagnostic, forensic, gene therapy and chromosome mapping procedures.
CC They are used to obtain upstream regulatory sequences and to design
CC expression and secretion vectors.
XX
SQ Sequence 325 BP; 100 A; 68 C; 68 G; 89 T; 0 other;

Query Match 92.0%; Score 18.4; DB 21; Length 325;
Best Local Similarity 95.0%; Pred. No. 37;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GATTAGCATATATAAATCTC 20
DB 39 GATTAGCATATATAAATCTC 20

RESULT 6
AAK61816
ID AAK61816 standard; cDNA; 645 BP.
XX
XX AAK61816;
XX
XX 06-NOV-2001 (first entry)
XX
DE Human immune/haematopoietic antigen encoding cDNA SEQ ID NO:6876.
XX
XX Human: Immune; haematopoietic; immune/haematopoietic antigen; cancer;
XX cytostatic; gene therapy; vaccine; metastasis; ss.
XX
XX Homo sapiens.
XX
XX WO200157182-A2.
XX
XX 09-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-US01354.
XX
XX 31-JAN-2000; 2000US-0179065.
XX
XX 04-FEB-2000; 2000US-0180628.
XX
XX 24-FEB-2000; 2000US-0184664.
XX
XX 02-MAR-2000; 2000US-0186350.
XX
XX 16-MAR-2000; 2000US-0189874.
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XX 17-MAR-2000; 2000US-0190076.
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XX 18-APR-2000; 2000US-0198123.
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XX 19-MAY-2000; 2000US-0205515.
XX
XX 07-JUN-2000; 2000US-0209467.
XX
XX 28-JUN-2000; 2000US-0214886.
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XX 30-JUN-2000; 2000US-0215135.
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XX 07-JUL-2000; 2000US-0216647.
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XX 07-JUL-2000; 2000US-0216880.
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XX 11-JUL-2000; 2000US-0217487.
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XX 11-JUL-2000; 2000US-0217496.
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XX 14-JUL-2000; 2000US-0218290.
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XX 26-JUL-2000; 2000US-0220963.
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XX 26-JUL-2000; 2000US-0220964.
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XX 14-AUG-2000; 2000US-0224518.
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XX 14-AUG-2000; 2000US-0224519.
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XX 14-AUG-2000; 2000US-0225213.
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XX 14-AUG-2000; 2000US-0225214.
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XX 14-AUG-2000; 2000US-0225266.
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XX 14-AUG-2000; 2000US-0225267.
XX
XX 14-AUG-2000; 2000US-0225268.
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XX 14-AUG-2000; 2000US-0225270.
XX
XX 14-AUG-2000; 2000US-0225447.
XX
XX 14-AUG-2000; 2000US-0225757.

PR 14-AUG-2000; 2000US-0225758.
PR 14-AUG-2000; 2000US-0225759.
PR 18-AUG-2000; 2000US-0226279.
PR 22-AUG-2000; 2000US-0226681.
PR 22-AUG-2000; 2000US-0226868.
PR 22-AUG-2000; 2000US-0227182.
PR 23-AUG-2000; 2000US-0227009.
PR 30-AUG-2000; 2000US-0228924.
PR 01-SEP-2000; 2000US-0229287.
PR 01-SEP-2000; 2000US-0229343.
PR 01-SEP-2000; 2000US-0229344.
PR 01-SEP-2000; 2000US-0229345.
PR 05-SEP-2000; 2000US-0229509.
PR 05-SEP-2000; 2000US-0229513.
PR 06-SEP-2000; 2000US-0230437.
PR 06-SEP-2000; 2000US-0230438.
PR 08-SEP-2000; 2000US-0231242.
PR 08-SEP-2000; 2000US-0231243.
PR 08-SEP-2000; 2000US-0231244.
PR 08-SEP-2000; 2000US-0231413.
PR 08-SEP-2000; 2000US-0231414.
PR 08-SEP-2000; 2000US-0232080.
PR 08-SEP-2000; 2000US-0232081.
PR 12-SEP-2000; 2000US-0231968.
PR 14-SEP-2000; 2000US-0232397.
PR 14-SEP-2000; 2000US-0232398.
PR 14-SEP-2000; 2000US-0232399.
PR 14-SEP-2000; 2000US-0232400.
PR 14-SEP-2000; 2000US-0232401.
PR 14-SEP-2000; 2000US-0233063.
PR 14-SEP-2000; 2000US-0233064.
PR 14-SEP-2000; 2000US-0233065.
PR 21-SEP-2000; 2000US-0234223.
PR 21-SEP-2000; 2000US-0234274.
PR 25-SEP-2000; 2000US-0234997.
PR 25-SEP-2000; 2000US-0234998.
PR 26-SEP-2000; 2000US-0235484.
PR 27-SEP-2000; 2000US-0235834.
PR 27-SEP-2000; 2000US-0235836.
PR 29-SEP-2000; 2000US-0236327.
PR 29-SEP-2000; 2000US-0236327.
PR 29-SEP-2000; 2000US-0236367.
PR 29-SEP-2000; 2000US-0236368.
PR 29-SEP-2000; 2000US-0236369.
PR 29-SEP-2000; 2000US-0236370.
PR 02-OCT-2000; 2000US-0236802.
PR 02-OCT-2000; 2000US-0237037.
PR 02-OCT-2000; 2000US-0237038.
PR 02-OCT-2000; 2000US-0237039.
PR 02-OCT-2000; 2000US-0237040.
PR 13-OCT-2000; 2000US-0239935.
PR 13-OCT-2000; 2000US-0239937.
PR 20-OCT-2000; 2000US-0240960.
PR 20-OCT-2000; 2000US-0241221.
PR 20-OCT-2000; 2000US-0241785.
PR 20-OCT-2000; 2000US-0241786.
PR 20-OCT-2000; 2000US-0241787.
PR 20-OCT-2000; 2000US-0241808.
PR 20-OCT-2000; 2000US-0241809.
PR 20-OCT-2000; 2000US-0241826.
PR 01-NOV-2000; 2000US-0244617.
PR 01-NOV-2000; 2000US-0244617.
PR 08-NOV-2000; 2000US-0246474.
PR 08-NOV-2000; 2000US-0246475.
PR 08-NOV-2000; 2000US-0246476.
PR 08-NOV-2000; 2000US-0246477.
PR 08-NOV-2000; 2000US-0246478.
PR 08-NOV-2000; 2000US-0246523.
PR 08-NOV-2000; 2000US-0246524.
PR 08-NOV-2000; 2000US-0246525.
PR 08-NOV-2000; 2000US-0246526.
PR 08-NOV-2000; 2000US-0246527.
PR 08-NOV-2000; 2000US-0246528.
PR 08-NOV-2000; 2000US-0246532.
PR 08-NOV-2000; 2000US-0246609.

PR 08-NOV-2000; 2000US-0246610.
PR 08-NOV-2000; 2000US-0246611.
PR 08-NOV-2000; 2000US-0246613.
PR 17-NOV-2000; 2000US-0249207.
PR 17-NOV-2000; 2000US-0249208.
PR 17-NOV-2000; 2000US-0249209.
PR 17-NOV-2000; 2000US-0249210.
PR 17-NOV-2000; 2000US-0249211.
PR 17-NOV-2000; 2000US-0249212.
PR 17-NOV-2000; 2000US-0249213.
PR 17-NOV-2000; 2000US-0249214.
PR 17-NOV-2000; 2000US-0249215.
PR 17-NOV-2000; 2000US-0249216.
PR 17-NOV-2000; 2000US-0249217.
PR 17-NOV-2000; 2000US-0249218.
PR 17-NOV-2000; 2000US-0249244.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249246.
PR 17-NOV-2000; 2000US-0249265.
PR 17-NOV-2000; 2000US-0249267.
PR 17-NOV-2000; 2000US-0249297.
PR 17-NOV-2000; 2000US-0249300.
PR 01-DEC-2000; 2000US-0250160.
PR 01-DEC-2000; 2000US-0250391.
PR 05-DEC-2000; 2000US-0251030.
PR 05-DEC-2000; 2000US-0251988.
PR 05-DEC-2000; 2000US-0256719.
PR 06-DEC-2000; 2000US-0251479.
PR 08-DEC-2000; 2000US-0251856.
PR 08-DEC-2000; 2000US-0251856.
PR 08-DEC-2000; 2000US-0251868.
PR 08-DEC-2000; 2000US-0251869.
PR 08-DEC-2000; 2000US-0251989.
PR 08-DEC-2000; 2000US-0251990.
PR 11-DEC-2000; 2000US-0254097.
PR 05-JAN-2001; 2001US-0253678.
XX
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
XX
PI Rosen CA, Barash SC, Ruben SM;
XX WPI, 2001-483426/52.
XX P-PSDB; AAM89035.
DR
XX
XX
PT Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
PT useful for preventing, diagnosing and/or treating cancers and
PT metastasis -
XX
XX
PS Claim 1; SEQ ID NO 6876; 3071pp + Sequence Listing; English.
XX
XX
CC AAK54951 to AAK64702 encode the human immune/hematopoietic antigen (I)
CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytosolic
CC activity, and can be used in gene therapy and vaccine production. (I)
CC proteins and polynucleotides may be used in the prevention, diagnosis and
CC treatment of diseases associated with inappropriate (I) expression. For
CC example, they may be used to treat disorders associated with decreased
CC expression by rectifying mutations or deletions in a patient's genome
CC that affect the activity of (I) by expressing inactive proteins or to
CC supplement the patient's own production of (I). Additionally, (I)
CC polynucleotides may be used to produce the secreted (I), by inserting
CC the nucleic acids into a host cell and culturing the cell to express the
CC protein. (I) proteins and polynucleotides may be used to prevent,
CC diagnose and treat immune/hematopoietic-related diseases, especially
CC cancers and cancer metastases of haematopoietic-derived cells. AAK64703
CC to AAK87694 represent human immune/hematopoietic antigen genomic
CC sequences from the present invention. AAK54942 to AAK54950 and AAM82169
CC represent sequences used in the exemplification of the present invention.
XX
SQ Sequence 645 BP; 193 A; 145 C; 118 G; 186 T; 3 other;

Query Match 92.0%; Score 18.4; DB 22; Length 645;
Best Local Similarity 95.0%; Pred. No. 37;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GATTGACATATAAATCTC 20
|||
Db 89 GAGTACGATATAAATCTC 108

RESULT 7
AAK81074
ID AAK81074 standard; DNA: 1972 BP.

XX AAK81074;

DT 07-NOV-2001 (first entry)

DE Human immune/haematopoietic antigen genomic sequence SEQ ID NO:35886.

KW Human; Immune; haematopoietic; immune/haematopoietic antigen; cancer;
KW cytostatic; gene therapy; vaccine; metastasis; ds.

XX Homo sapiens.

PM WO200157182-A2.

PD 09-AUG-2001.

PF 17-JAN-2001; 2001WO-US01354.

XX 31-JAN-2000; 2000US-0179065.

PR 04-FEB-2000; 2000US-0180628.

PR 24-FEB-2000; 2000US-0184664.

PR 02-MAR-2000; 2000US-0186350.

PR 16-MAR-2000; 2000US-0189874.

PR 17-MAR-2000; 2000US-0190076.

PR 18-APR-2000; 2000US-0198123.

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PR 07-JUL-2000; 2000US-0216880.

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PR 14-JUL-2000; 2000US-0218290.

PR 26-JUL-2000; 2000US-0220963.

PR 14-AUG-2000; 2000US-0220964.

PR 14-AUG-2000; 2000US-0224518.

PR 14-AUG-2000; 2000US-0224519.

PR 14-AUG-2000; 2000US-0225213.

PR 14-AUG-2000; 2000US-0225214.

PR 14-AUG-2000; 2000US-0225266.

PR 14-AUG-2000; 2000US-0225267.

PR 14-AUG-2000; 2000US-0225268.

PR 14-AUG-2000; 2000US-0225270.

PR 14-AUG-2000; 2000US-0225447.

PR 14-AUG-2000; 2000US-0225757.

PR 14-AUG-2000; 2000US-0225758.

PR 14-AUG-2000; 2000US-0225759.

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PR 22-AUG-2000; 2000US-0226866.

PR 23-AUG-2000; 2000US-0227182.

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PR 30-AUG-2000; 2000US-0228924.

PR 01-SEP-2000; 2000US-0229287.

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PR 01-SEP-2000; 2000US-0229344.

PR 01-SEP-2000; 2000US-0229345.

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PR 05-SEP-2000; 2000US-0229513.

PR 06-SEP-2000; 2000US-0230437.

PR 06-SEP-2000; 2000US-0230438.

PR 08-SEP-2000; 2000US-0231242.

PR 08-SEP-2000; 2000US-0231243.

PR 08-SEP-2000; 2000US-0231244.

PR 08-SEP-2000; 2000US-0231413.

PR 08-SEP-2000; 2000US-0231414.

PR 08-SEP-2000; 2000US-0232080.

PR 08-SEP-2000; 2000US-0232081.

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PR 14-SEP-2000; 2000US-0232399.

PR 14-SEP-2000; 2000US-0232400.

PR 14-SEP-2000; 2000US-0232401.

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PR 21-SEP-2000; 2000US-0234274.

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PR 25-SEP-2000; 2000US-0234998.

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PR 27-SEP-2000; 2000US-0235835.

PR 29-SEP-2000; 2000US-0236327.

PR 29-SEP-2000; 2000US-0236367.

PR 29-SEP-2000; 2000US-0236368.

PR 29-SEP-2000; 2000US-0236369.

PR 29-SEP-2000; 2000US-0236370.

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PR 13-OCT-2000; 2000US-0239937.

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PR 20-OCT-2000; 2000US-0241787.

PR 20-OCT-2000; 2000US-0241808.

PR 20-OCT-2000; 2000US-0241809.

PR 20-OCT-2000; 2000US-0241826.

PR 01-NOV-2000; 2000US-0244617.

PR 08-NOV-2000; 2000US-0246474.

PR 08-NOV-2000; 2000US-0246475.

PR 08-NOV-2000; 2000US-0246476.

PR 08-NOV-2000; 2000US-0246477.

PR 08-NOV-2000; 2000US-0246478.

PR 08-NOV-2000; 2000US-0246523.

PR 08-NOV-2000; 2000US-0246524.

PR 08-NOV-2000; 2000US-0246525.

PR 08-NOV-2000; 2000US-0246526.

PR 08-NOV-2000; 2000US-0246527.

PR 08-NOV-2000; 2000US-0246528.

PR 08-NOV-2000; 2000US-0246532.

PR 08-NOV-2000; 2000US-0246609.

PR 08-NOV-2000; 2000US-0246610.

PR 08-NOV-2000; 2000US-0246611.

PR 08-NOV-2000; 2000US-0246613.

PR 17-NOV-2000; 2000US-0249207.

PR 17-NOV-2000; 2000US-0249208.

PR 17-NOV-2000; 2000US-0249209.

PR 17-NOV-2000; 2000US-0249210.

PR 17-NOV-2000; 2000US-0249211.

PR 17-NOV-2000; 2000US-0249212.

PR 17-NOV-2000; 2000US-0249213.

PR 17-NOV-2000; 2000US-0249215.

PR 17-NOV-2000; 2000US-0249216.

PR 17-NOV-2000; 2000US-0249217.

PR 17-NOV-2000; 2000US-0249218.

PR 17-NOV-2000; 2000US-0249244.

PR 17-NOV-2000; 2000US-0249245.

PR 17-NOV-2000; 2000US-0249264.

PR 17-NOV-2000; 2000US-0249265.

[illegible]

PR 11-JUL-2000; 2000US-0217496.
PR 14-JUL-2000; 2000US-0218290.
PR 26-JUL-2000; 2000US-0220963.
PR 26-JUL-2000; 2000US-0220964.
PR 14-AUG-2000; 2000US-0224518.
PR 14-AUG-2000; 2000US-0224519.
PR 14-AUG-2000; 2000US-0225211.
PR 14-AUG-2000; 2000US-0225214.
PR 14-AUG-2000; 2000US-0225266.
PR 14-AUG-2000; 2000US-0225267.
PR 14-AUG-2000; 2000US-0225268.
PR 14-AUG-2000; 2000US-0225270.
PR 14-AUG-2000; 2000US-0225447.
PR 14-AUG-2000; 2000US-0225757.
PR 14-AUG-2000; 2000US-0225758.
PR 14-AUG-2000; 2000US-0225759.
PR 18-AUG-2000; 2000US-0226279.
PR 22-AUG-2000; 2000US-0226681.
PR 22-AUG-2000; 2000US-0226868.
PR 22-AUG-2000; 2000US-0227182.
PR 23-AUG-2000; 2000US-0227009.
PR 30-AUG-2000; 2000US-0228924.
PR 01-SEP-2000; 2000US-0229287.
PR 01-SEP-2000; 2000US-0229343.
PR 01-SEP-2000; 2000US-0229344.
PR 01-SEP-2000; 2000US-0229345.
PR 05-SEP-2000; 2000US-0229509.
PR 05-SEP-2000; 2000US-0229513.
PR 06-SEP-2000; 2000US-0230437.
PR 06-SEP-2000; 2000US-0230438.
PR 08-SEP-2000; 2000US-0231242.
PR 08-SEP-2000; 2000US-0231243.
PR 08-SEP-2000; 2000US-0231244.
PR 08-SEP-2000; 2000US-0231413.
PR 08-SEP-2000; 2000US-0231414.
PR 08-SEP-2000; 2000US-0232080.
PR 08-SEP-2000; 2000US-0232081.
PR 12-SEP-2000; 2000US-0231968.
PR 14-SEP-2000; 2000US-0233397.
PR 14-SEP-2000; 2000US-0233398.
PR 14-SEP-2000; 2000US-0233399.
PR 14-SEP-2000; 2000US-0233400.
PR 14-SEP-2000; 2000US-0233401.
PR 14-SEP-2000; 2000US-0233063.
PR 14-SEP-2000; 2000US-0233064.
PR 14-SEP-2000; 2000US-0233065.
PR 21-SEP-2000; 2000US-0234223.
PR 21-SEP-2000; 2000US-0234274.
PR 25-SEP-2000; 2000US-0234997.
PR 25-SEP-2000; 2000US-0234998.
PR 26-SEP-2000; 2000US-0234984.
PR 27-SEP-2000; 2000US-0235834.
PR 27-SEP-2000; 2000US-0235835.
PR 29-SEP-2000; 2000US-0236327.
PR 29-SEP-2000; 2000US-0236367.
PR 29-SEP-2000; 2000US-0236368.
PR 29-SEP-2000; 2000US-0236369.
PR 29-SEP-2000; 2000US-0236370.
PR 02-OCT-2000; 2000US-0236802.
PR 02-OCT-2000; 2000US-0237037.
PR 02-OCT-2000; 2000US-0237038.
PR 02-OCT-2000; 2000US-0237039.
PR 02-OCT-2000; 2000US-0237040.
PR 13-OCT-2000; 2000US-0239935.
PR 13-OCT-2000; 2000US-0239937.
PR 20-OCT-2000; 2000US-0240960.
PR 20-OCT-2000; 2000US-0241221.
PR 20-OCT-2000; 2000US-0241785.
PR 20-OCT-2000; 2000US-0241786.
PR 20-OCT-2000; 2000US-0241787.
PR 20-OCT-2000; 2000US-0241808.
PR 20-OCT-2000; 2000US-0241809.
PR 20-OCT-2000; 2000US-0241826.
PR 20-OCT-2000; 2000US-0241826.

PR 01-NOV-2000; 2000US-0244617.
PR 08-NOV-2000; 2000US-0246474.
PR 08-NOV-2000; 2000US-0246475.
PR 08-NOV-2000; 2000US-0246476.
PR 08-NOV-2000; 2000US-0246477.
PR 08-NOV-2000; 2000US-0246478.
PR 08-NOV-2000; 2000US-0246523.
PR 08-NOV-2000; 2000US-0246524.
PR 08-NOV-2000; 2000US-0246525.
PR 08-NOV-2000; 2000US-0246526.
PR 08-NOV-2000; 2000US-0246527.
PR 08-NOV-2000; 2000US-0246528.
PR 08-NOV-2000; 2000US-0246532.
PR 08-NOV-2000; 2000US-0246609.
PR 08-NOV-2000; 2000US-0246610.
PR 08-NOV-2000; 2000US-0246613.
PR 17-NOV-2000; 2000US-0249207.
PR 17-NOV-2000; 2000US-0249208.
PR 17-NOV-2000; 2000US-0249209.
PR 17-NOV-2000; 2000US-0249210.
PR 17-NOV-2000; 2000US-0249211.
PR 17-NOV-2000; 2000US-0249212.
PR 17-NOV-2000; 2000US-0249213.
PR 17-NOV-2000; 2000US-0249214.
PR 17-NOV-2000; 2000US-0249215.
PR 17-NOV-2000; 2000US-0249216.
PR 17-NOV-2000; 2000US-0249217.
PR 17-NOV-2000; 2000US-0249218.
PR 17-NOV-2000; 2000US-0249244.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249246.
PR 17-NOV-2000; 2000US-0249264.
PR 17-NOV-2000; 2000US-0249265.
PR 17-NOV-2000; 2000US-0249266.
PR 17-NOV-2000; 2000US-0249267.
PR 17-NOV-2000; 2000US-0249268.
PR 17-NOV-2000; 2000US-0249300.
PR 01-DEC-2000; 2000US-0250160.
PR 01-DEC-2000; 2000US-0250391.
PR 05-DEC-2000; 2000US-0251030.
PR 05-DEC-2000; 2000US-0251031.
PR 05-DEC-2000; 2000US-0251988.
PR 05-DEC-2000; 2000US-0256719.
PR 06-DEC-2000; 2000US-0251479.
PR 06-DEC-2000; 2000US-0251856.
PR 08-DEC-2000; 2000US-0251868.
PR 08-DEC-2000; 2000US-0251869.
PR 08-DEC-2000; 2000US-0251989.
PR 08-DEC-2000; 2000US-0251990.
PR 11-DEC-2000; 2000US-0254097.
PR 05-JAN-2001; 2001US-0259678.
PR (HUMA-) HUMAN GENOME SCI INC.
PR PA
PR XX
PR XX
PR PI
PR WPI: 2001-483426/52.
PR DR
PR XX
PR PT
PR PT
PR PT
PR PS
PR XX
CC AAK54951 to AAK64702 encode the human Immune/Hematopoietic antigen (I)
CC amino acid sequences given in AAK82170 to AAK91921. (I) have cytosolic
CC activity, and can be used in gene therapy and vaccine production. (I)
CC proteins and polynucleotides may be used in the prevention, diagnosis and
CC treatment of diseases associated with inappropriate (I) expression. For
CC example, they may be used to treat disorders associated with decreased
CC expression by rectifying mutations or deletions in a patient's genome
CC that affect the activity of (I) by expressing inactive proteins or to
CC supplement the patients own production of (I). Additionally, (I)
CC polynucleotides may be used to produce the secreted (I), by inserting
CC the nucleic acids into a host cell and culturing the cell to express the

CC	protein. (1) proteins and polynucleotides may be used to prevent,
CC	diagnose and treat immune/haematopoietic-related diseases, especially
CC	cancers and cancer metastases of haematopoietic-derived cells. AAK64703
CC	to AAK67694 represent human immune/haematopoietic antigen genomic
CC	sequences from the present invention. AAK54942 to AAK54950 and AAM82169
CC	represent sequences used in the exemplification of the present invention.
XX	
XX	Sequence 1976 BP; 663 A; 409 C; 350 G; 554 T; 0 other;
SO	
	Query Match 92.0%; Score 18.4; DB 22; Length 1976;
	Best Local Similarity 95.0%; Pred. No. 37;
	Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0
OY	1 GATTAGCATTAATAATCTC 20
Db	89 GAGTAGCATTAATAATCTC 108
RESULT 10	
ACCS9841/C	
ID	ACCS9841 standard; DNA; 3153 BP.
AC	
ACCS9841;	
DT	26-JAN-2001 (first entry)
XX	
DE	Human secreted protein encoding DNA clone vo31 1.
KW	Secreted protein; human; autoimmune disorder; multiple sclerosis; ulcer;
KW	systemic lupus erythematosus; rheumatoid arthritis; anaemia; stroke;
KW	hematopoiesis regulation; tissue regrowth; wound healing; haemophilia;
KW	Alzheimer's disease; Parkinson's disease; Shy-draeger syndrome; cancer;
KW	contraceptive; infection; growth inhibition; hyperproliferative disorder;
KW	psoriasis; ds.
XX	
OS	Homo sapiens.
XX	
PN	WO20005375-A1.
PD	21-SEP-2000.
XX	
PF	17-MAR-2000; 2000MO-US07285.
XX	
PR	17-MAR-1999; 99US-0124808.
PR	17-MAR-1999; 99US-0124916.
PR	17-AUG-1999; 99US-0148639.
PR	01-OCT-1998; 99US-0157247.
PR	29-NOV-1999; 99US-0167824.
PR	15-FEB-2000; 2000US-0182711.
XX	
PA	(ALPH-) ALPHAGENE INC.
PI	
PI	Valenzuela D, Yuan O, Hoffman H, Hall J, Rapiejko P;
DR	WPI: 2000-638211/61.
DR	P-PSDB: AAB34740.
PT	
PT	Novel proteins and polypeptides useful for the treatment of e.g
PT	multiple sclerosis, systemic lupus erythematosus, rheumatoid arthritis,
PT	cancer, Alzheimer's disease, Parkinson's disease, stroke, anemia and
PT	ulcers -
XX	
PS	Claim 116; Page 455-456; 493pp; English.
XX	
CC	This invention relates to 59 human secreted proteins and the nucleotide
CC	sequences encoding them. Sequences AAC59788-C59846 and AAB34687-B34745
CC	represent the proteins and their encoding nucleotide sequences, and
CC	sequences AAB34746-B34771 represent fragments of the proteins. Probes
CC	for the DNA sequences are represented by sequences AAC59847-C59956. The
CC	proteins exhibit neuroprotective, dermatological, immunosuppressive,
CC	antiinflammatory, antianaemic, nootropic, antiparkinsonian,
CC	cerebroprotective, haemostatic, vulnerary, cytostatic, antiporiatic,
CC	antibacterial, virucide, and fungicide activity. The proteins and
CC	

CC	nucleotide sequences are useful as nutritional sources or supplements
CC	and in research. The proteins are useful for treating immune deficiency
CC	and disorders, which may be genetic or resulting from infections,
CC	autoimmune disorders such as multiple sclerosis, systemic lupus
CC	erythematosus, rheumatoid arthritis, and for treating myeloid or lymphoid
CC	cell deficiencies such as anaemias by regulating haematopoiesis. The
CC	proteins are also useful in compositions for bone, cartilage, tendon,
CC	ligament and/or nerve tissue growth or regeneration, for wound healing,
CC	tissue repair and replacement and in the treatment of wounds, incisions
CC	and ulcers. Other uses include in the treatment of central and
CC	peripheral nervous system and neuropathies such as Alzheimer's and
CC	Parkinson's diseases and Shy-Drager syndrome, and mechanical and
CC	traumatic disorders, such as spinal cord disorders, head trauma and
CC	stroke. The proteins may also be used as a contraceptive, and for
CC	treating coagulation disorders such as haemophilias. The protein and
CC	nucleotide sequences with cadherin activity are useful for treating
CC	cancer. Other uses for the protein include for inhibiting the growth,
CC	infection or function of, or killing, infectious agents such as bacteria,
CC	virus, fungi and other parasites, for effecting bodily characteristics
CC	such as height, weight, hair colour, effecting biorhythms or cardiac
CC	cycles or rhythms, effecting metabolism, catabolism, anabolism,
CC	processing, utilization, storage or elimination of dietary fat, lipid,
CC	protein, carbohydrate, vitamins, minerals, cofactors, effecting
CC	behavioural characteristics, providing analgesic effects and for treating
CC	hyperproliferative disorders such as psoriasis.
CC	
SQ	Sequence 3153 BP; 938 A; 709 C; 647 G; 859 T; 0 other;
Query Match	87.0%; Score 17.4; DB 21; Length 3153;
Best Local Similarity	94.7%; Pred. No. 1e+02;
Matches 18; Conservative	0; Mismatches 1; Indels 0; Gaps 0;
OY	2 ATTACGATATATAATCTC 20
Db	1944 AFTGCATATATAATCTC 1926
RESULT 11	
ID	AAK95332/c
XX	AAK95332 standard; DNA; 401 BP.
AC	
XX	AAK95332;
DT	17-DEC-2001 (first entry)
DE	
XX	Human neuregulin gene single nucleotide polymorphism SNP8NRC6787.
XX	
KW	Human; neuregulin-1 associated gene 1; NR1AG1; Schizophrenia gene;
RW	gene therapy; single nucleotide polymorphism; SNP; ds.
OS	Homo sapiens.
XX	
XX	WO200164876-A2.
PN	
XX	07-SEP-2001.
PD	
PF	28-FEB-2001; 2001WO-US06376.
XX	
XX	28-FEB-2000; 2000US-0515715.
XX	
PA	(DECO-) DECODE GENETICS EHF.
XX	
PI	Stefansson H, Steinthorsdottir V, Gulcher JR;
DR	WPI; 2001-550179/61.
XX	
PT	Neuregulin-1 associated gene 1 nucleic acids and fragments, useful for
XX	preventing diagnosing and treating schizophrenia -
XX	Disclosure; Page 510; 750pp; English.
CC	
CC	This sequence represents a single nucleotide polymorphism (SNP) of the
CC	human neuregulin-1 associated gene 1 (NR1AG1) of the invention. The

CC NR1AG1 gene is also referred to as the human Schizophrenia gene. The
CC invention also relates to fragments or variants of the gene and the
CC NR1AG1 polypeptides they encode. The NR1AG1 nucleic acids and
CC polypeptides may be used in the prevention, diagnosis and treatment of
CC diseases associated with inappropriate NR1AG1 expression. For example,
CC they may be used to treat disorders associated with decreased expression
CC by rectifying mutations or deletions in a patient's genome that affect
CC the activity of NR1AG1 by expressing inactive proteins or to supplement
CC the patients own production of NR1AG1. Additionally, the gene may be
CC used to produce NR1AG1 polypeptides, by inserting the nucleic acids into
CC a host cell and culturing the cell to express the protein. The gene may
CC also be used as DNA probes and primers in diagnostic assays to detect and
CC quantitate the presence of similar nucleic acids in samples, and
CC therefore which patients may be in need of restorative therapy. The
CC NR1AG1 polypeptides may also be used as antigens in the production of
CC antibodies against NR1AG1 and in assays to identify modulators of
CC NR1AG1 expression and activity. Anti-NR1AG1 antibodies and antagonists
CC may also be used to down regulate expression and activity. Anti-NR1AG1
CC antibodies may also be used as diagnostic agents for detecting the
CC presence of NR1AG1 polypeptides in samples. NR1AG1 is associated with
CC schizophrenia which may be prevented, diagnosed and/or treated by the
CC above methods.

XX Sequence 401 BP; 119 A; 96 C; 84 G; 98 T; 4 other;

Query Match 82.0%; Score 16.4; DB 22; Length 401;

Best Local Similarity 94.4%; Pred. No. 2.8e+02;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GATTAGCATATTAATTC 18

||||| |||||||||

DB 358 GATTTCATATTAATTC 341

RESULT 12

AAK95333/C

ID AAK95333 standard; DNA; 401 BP.

XX AAK95333;

DT 17-DEC-2001 (first entry)

DE Human neuregulin gene single nucleotide polymorphism SNP8NRG6844.

XX Human; neuregulin-1 associated gene 1; NR1AG1; Schizophrenia gene;

KW gene therapy; single nucleotide polymorphism; SNP; ds.

XX Homo sapiens.

OS

PN WO200164876-A2.

XX 07-SEP-2001.

PD 28-FEB-2001; 2001WO-US06376.

XX 28-FEB-2000; 2000US-0515715.

PR (DECO-) DECODE GENETICS EHF.

XX (DECO-) DECODE GENETICS EHF.

PI Stefansson H, Steinthorsdottir V, Gulcher JR;

XX WPI; 2001-550179/61.

DR Neuregulin-1 associated gene 1 nucleic acids and fragments, useful for

XX preventing diagnosing and treating schizophrenia -

PT Disclosure; Page 510; 750pp; English.

XX This sequence represents a single nucleotide polymorphism (SNP) of the

CC human neuregulin-1 associated gene 1 (NR1AG1) of the invention. The

CC NR1AG1 gene is also referred to as the human Schizophrenia gene. The

CC invention also relates to fragments or variants of the gene and the

CC NR1AG1 polypeptides they encode. The NR1AG1 nucleic acids and

CC polypeptides may be used in the prevention, diagnosis and treatment of
CC diseases associated with inappropriate NR1AG1 expression. For example,
CC they may be used to treat disorders associated with decreased expression
CC by rectifying mutations or deletions in a patient's genome that affect
CC the activity of NR1AG1 by expressing inactive proteins or to supplement
CC the patients own production of NR1AG1. Additionally, the gene may be
CC used to produce NR1AG1 polypeptides, by inserting the nucleic acids into
CC a host cell and culturing the cell to express the protein. The gene may
CC also be used as DNA probes and primers in diagnostic assays to detect and
CC quantitate the presence of similar nucleic acids in samples, and
CC therefore which patients may be in need of restorative therapy. The
CC NR1AG1 polypeptides may also be used as antigens in the production of
CC antibodies against NR1AG1 and in assays to identify modulators of
CC NR1AG1 expression and activity. Anti-NR1AG1 antibodies and antagonists
CC may also be used to down regulate expression and activity. Anti-NR1AG1
CC antibodies may also be used as diagnostic agents for detecting the
CC presence of NR1AG1 polypeptides in samples. NR1AG1 is associated with
CC schizophrenia which may be prevented, diagnosed and/or treated by the
CC above methods.

XX Sequence 401 BP; 114 A; 89 C; 92 G; 103 T; 3 other;

Query Match 82.0%; Score 16.4; DB 22; Length 401;

Best Local Similarity 94.4%; Pred. No. 2.8e+02;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GATTAGCATATTAATTC 18

||||| |||||||||

DB 301 GATTTCATATTAATTC 284

RESULT 13

AAK96579/C

ID AAK96579 standard; DNA; 401 BP.

XX AAK96579;

DT 17-DEC-2001 (first entry)

DE Human neuregulin gene insertion/deletion DNP8NRG1.

XX Human; neuregulin-1 associated gene 1; NR1AG1; Schizophrenia gene;

KW gene therapy; insertion; deletion; ds.

XX Homo sapiens.

OS

PN WO200164876-A2.

XX 07-SEP-2001.

PD 28-FEB-2001; 2001WO-US06376.

XX 28-FEB-2000; 2000US-0515715.

PR (DECO-) DECODE GENETICS EHF.

XX (DECO-) DECODE GENETICS EHF.

PI Stefansson H, Steinthorsdottir V, Gulcher JR;

XX WPI; 2001-550179/61.

DR Neuregulin-1 associated gene 1 nucleic acids and fragments, useful for

XX preventing diagnosing and treating schizophrenia -

PT Disclosure; Page 729; 750pp; English.

XX This sequence represents an insertion/deletion variant of the human

CC neuregulin-1 associated gene 1 (NR1AG1) of the invention. The NR1AG1

CC gene is also referred to as the human Schizophrenia gene. The invention

CC also relates to fragments or variants of the gene and the NR1AG1

CC polypeptides they encode. The NR1AG1 nucleic acids and polypeptides may

CC be used in the prevention, diagnosis and treatment of diseases associated

CC with inappropriate NR1AG1 expression. For example, they may be used to

CC treat disorders associated with decreased expression by rectifying

CC mutations or deletions in a patient's genome that affect the activity of
CC NRGIAG1 by expressing inactive proteins or to supplement the patients own
CC production of NRGIAG1. Additionally, the gene may be used to produce
CC NRGIAG1 polypeptides, by inserting the nucleic acids into a host cell
CC and culturing the cell to express the protein. The gene may also be used
CC as DNA probes and primers in diagnostic assays to detect and quantitate
CC the presence of similar nucleic acids in samples, and therefore which
CC patients may be in need of restorative therapy. The NRGIAG1 polypeptides
CC may also be used as antigens in the production of antibodies against
CC NRGIAG1 and in assays to identify modulators of NRGIAG1 expression and
CC activity. Anti-NRGIAG1 antibodies and antagonists may also be used to
CC down regulate expression and activity. Anti-NRGIAG1 antibodies may
CC also be used as diagnostic agents for detecting the presence of NRGIAG1
CC polypeptides in samples. NRGIAG1 is associated with schizophrenia which
CC may be prevented, diagnosed and/or treated by the above methods.

SO Sequence 401 BP; 124 A; 78 C; 80 G; 117 T; 2 other;

Query Match 82.0%; Score 16.4; DB 22; Length 401;
Best Local Similarity 94.4%; Pred. No. 2.8e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GATTACCATATAAATC 18
DB 186 GATTTCATATAAATC 169

RESULT 14
AAK96581/C
ID AAK96581 standard; DNA; 401 BP.

AC AAK96581;

DT 17-DEC-2001 (first entry)

DE Human neuregulin gene insertion/deletion DNP8NRG3.

KW Human; neuregulin-1 associated gene 1; NRGIAG1; Schizophrenia gene;

KM gene therapy; Insertion; deletion; ds.

OS Homo sapiens.

PN WO200164876-A2.

PD 07-SEP-2001.

PF 28-FEB-2001; 2001WO-US06376.

PR 28-FEB-2000; 2000US-0515715.

PA (DECO-) DECODE GENETICS EHF.

PI Stefansson H, Steinthorsdottir V, Gulcher JR;

DR WPI; 2001-550179/61.

PT Neuregulin-1 associated gene 1 nucleic acids and fragments, useful for
preventing diagnosing and treating schizophrenia

PS Disclosure: Page 729; 750pp; English.

CC This sequence represents an insertion/deletion variant of the human
CC neuregulin-1 associated gene 1 (NRGIAG1) of the invention. The NRGIAG1
CC gene is also referred to as the human Schizophrenia gene. The invention
CC also relates to fragments or variants of the gene and the NRGIAG1
CC polypeptides they encode. The NRGIAG1 nucleic acids and polypeptides may
CC be used in the prevention, diagnosis and treatment of diseases associated
CC with inappropriate NRGIAG1 expression. For example, they may be used to
CC treat disorders associated with decreased expression by rectifying
CC mutations or deletions in a patient's genome that affect the activity of
CC NRGIAG1 by expressing inactive proteins or to supplement the patients own
CC production of NRGIAG1. Additionally, the gene may be used to produce
CC NRGIAG1 polypeptides, by inserting the nucleic acids into a host cell

CC and culturing the cell to express the protein. The gene may also be used
CC as DNA probes and primers in diagnostic assays to detect and quantitate
CC the presence of similar nucleic acids in samples, and therefore which
CC patients may be in need of restorative therapy. The NRGIAG1 polypeptides
CC may also be used as antigens in the production of antibodies against
CC NRGIAG1 and in assays to identify modulators of NRGIAG1 expression and
CC activity. Anti-NRGIAG1 antibodies and antagonists may also be used to
CC down regulate expression and activity. Anti-NRGIAG1 antibodies may
CC also be used as diagnostic agents for detecting the presence of NRGIAG1
CC polypeptides in samples. NRGIAG1 is associated with schizophrenia which
CC may be prevented, diagnosed and/or treated by the above methods.

SO Sequence 401 BP; 119 A; 74 C; 85 G; 123 T; 0 other;

Query Match 82.0%; Score 16.4; DB 22; Length 401;
Best Local Similarity 94.4%; Pred. No. 2.8e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GATTACCATATAAATC 18
DB 90 GATTTCATATAAATC 73

RESULT 15
AAK96825/C
ID AAK96825 standard; DNA; 401 BP.

AC AAK96825;

DT 17-DEC-2001 (first entry)

DE Human neuregulin gene single nucleotide polymorphism SNP8NRG6787.

KW Human; neuregulin 1 gene; schizophrenia; gene therapy; SNP;

KM single nucleotide polymorphism; ds.

OS Homo sapiens.

PN WO200164877-A2.

PD 07-SEP-2001.

PF 28-FEB-2001; 2001WO-US06377.

PR 28-FEB-2000; 2000US-0515716.

PA (DECO-) DECODE GENETICS EHF.

PI Stefansson H, Steinthorsdottir V, Gulcher JR;

DR WPI; 2001-514841/56.

PT Neuregulin 1 nucleic acids and proteins useful for diagnosing
preventing and treating schizophrenia

PS Disclosure: Page 95; 756pp; English.

CC This sequence represents a single nucleotide polymorphism (SNP)
CC from the human neuregulin 1 gene of the invention.
CC The invention also relates to fragments or variants of the neuregulin 1
CC gene. The gene and its proteins may be used in the prevention, diagnosis
CC and treatment of diseases associated with inappropriate neuregulin 1
CC expression, such as schizophrenia. For example they may be used to treat
CC disorders associated with decreased neuregulin 1 expression by rectifying
CC mutations or deletions in a patient's genome that affect the activity of
CC neuregulin 1 by expressing inactive proteins or to supplement the
CC patients own production of polypeptides. Additionally, the gene may be
CC used to produce the neuregulin 1 protein, by inserting the nucleic acids
CC into a host cell and culturing the cell to express the protein. The gene
CC and its complementary sequences may also be used as DNA probes in
CC diagnostic assays to detect and quantitate the presence of similar
CC nucleic acids in samples, and therefore which patients may be in need of
CC restorative therapy. The protein may also be used as antigens in the

CC production of antibodies against neuregulin 1 and in assays to identify
 CC modulators of neuregulin 1 expression and activity. The antibodies and
 CC antagonists may also be used to down regulate expression and activity.
 CC The antibodies may also be used as diagnostic agents for detecting the
 CC presence of neuregulin 1 in samples.

XX
 SQ Sequence 401 BP; 119 A; 96 C; 84 G; 98 T; 4 other;

Query Match 82.0%; Score 16.4; DB 22; Length 401;
 Best Local Similarity 94.4%; Pred. No. 2.8e+02;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GATTAGCATATTAATC 18
 |||| |||||
 DB 358 GATTGACATATTAATC 341

Search completed: October 6, 2002, 15:21:06
 Job time : 290 secs

GenCore version 5.1.3
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OM nucleic - nucleic search, using SW model

Run on: October 6, 2002, 13:13:14 ; Search time 2261 Seconds
(without alignments)
185.109 Million cell updates/sec

Title: US-09-754-468-47
Perfect score: 20
Sequence: 1 gattagcataataaattctc 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues
Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 08
Maximum Match 1008
Listing first 45 summaries

Database : GenEmbl:
1: gb.ba:*
2: gb.htg:*
3: gb.in:*
4: gb.om:*
5: gb.ov:*
6: gb.pat:*
7: gb.ph:*
8: gb.pl:*
9: gb.pr:*
10: gb.ro:*
11: gb.sts:*
12: gb.sy:*
13: gb.un:*
14: gb.vi:*
15: em.ba:*
16: em.fun:*
17: em.hum:*
18: em.in:*
19: em.mu:*
20: em.om:*
21: em.or:*
22: em.ov:*
23: em.pat:*
24: em.ph:*
25: em.pl:*
26: em.ro:*
27: em.sts:*
28: em.un:*
29: em.vi:*
30: em.htg.hum:*
31: em.htg.inv:*
32: em.htg.other:*
33: em.htg.inh:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No. Score Query Match Length DB ID Description

1	20	100.0	20	6	AX191765	AX191765 Sequence
2	20	100.0	836	6	ARI49152	ARI49152 Sequence
3	20	100.0	836	6	ARI49153	ARI49153 Sequence
4	20	100.0	2048	1	ECOEENVA	M19211 E.coli cell
5	20	100.0	2754	6	E27365	E27365 Treponema P
6	20	100.0	3811	1	ECOSECA	M20791 Escherichia
7	20	100.0	12434	1	AE000119	AE000119 Escherich
8	20	100.0	12518	1	AE005186	AE005186 Escherich
9	20	100.0	28277	1	EC2MIN	X55034 E. coli 2 m
10	20	100.0	28277	6	AX191720	AX191720 Sequence
11	20	100.0	111401	1	EC0110K	D10483 E.coli K12
12	19	95.0	155976	2	AC055113	AC055113 Homo sapi
13	19	95.0	179212	9	AC084877	AC084877 Homo sapi
14	18.4	92.0	22286	1	AE008700	AE008700 Salmonell
15	18.4	92.0	127603	9	AC036102	AC036102 Homo sapi
16	18.4	92.0	138846	2	AC068478	AC068478 Homo sapi
17	18.4	92.0	156978	2	AC023379	AC023379 Homo sapi
18	18.4	92.0	157385	9	AC006022	AC006022 Homo sapi
19	18.4	92.0	165102	2	AC074279	AC074279 Homo sapi
20	18.4	92.0	166427	2	AC069593	AC069593 Homo sapi
21	18.4	92.0	168004	2	AC019203	AC019203 Homo sapi
22	18.4	92.0	172579	9	AC008064	AC008064 Homo sapi
23	18.4	92.0	183084	2	AC025936	AC025936 Homo sapi
24	18.4	92.0	188049	2	AC069391	AC069391 Homo sapi
25	18.4	92.0	193588	2	AC025370	AC025370 Homo sapi
26	18.4	92.0	194782	2	AC092433	AC092433 Homo sapi
27	18.4	92.0	200475	9	AC079271	AC079271 Homo sapi
28	18.4	92.0	201629	2	AL158047	AL158047 Human DNA
29	18.4	92.0	204790	2	AC013452	AC013452 Homo sapi
30	18.4	92.0	251050	1	AF627265	AF627265 Salmonell
31	18.4	92.0	347253	9	AF363578	AF363578 Homo sapi
32	18	90.0	51869	2	AC094868	AC094868 Rattus no
33	18	90.0	175456	2	AL359976	AL359976 Homo sapi
34	18	90.0	195290	2	AC074364	AC074364 Homo sapi
35	17.4	87.0	942	33	AC038729	AC038729 Giardia 1
36	17.4	87.0	63769	2	AC079334	AC079334 Homo sapi
37	17.4	87.0	73610	2	AL359735	AL359735 Human DNA
38	17.4	87.0	102990	2	AP000801	AP000801 Homo sapi
39	17.4	87.0	109568	2	AP000840	AP000840 Homo sapi
40	17.4	87.0	126755	9	HSBJ18C9	AL1049709 Human DNA
41	17.4	87.0	138969	9	AC060771	AC060771 Homo sapi
42	17.4	87.0	142278	2	AC005177	AC005177 Homo sapi
43	17.4	87.0	150406	2	AC098684	AC098684 Mus muscu
44	17.4	87.0	161742	2	AC036204	AC036204 Homo sapi
45	17.4	87.0	161742	2	AC036204	AC036204 Homo sapi

ALIGNMENTS

RESULT 1
AX191765
LOCUS AX191765 20 bp DNA linear PAT 15-AUG-2001
DEFINITION Sequence 47 from Patent WO0149775.
ACCESSION AX191765
VERSION AX191765.1 GI:15209934
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 20)
AUTHORS Iversen, P.L.
TITLE Antisense antibacterial cell division composition and method
JOURNAL Patent: WO 0149775-A 47 12-JUL-2001;
Avi Biopharma, Inc. (US)
FEATURES
source location/Qualifiers
1..20
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Oligonucleotide"

100 LATE
SAMP ENV

	Query Match	100.0%;	Score 20;	DB 6;	Length 20;
	Best Local Similarity	100.0%;	Pred. No. 2.*e+02;		
	Matches	20;	Conservative 0;	Mismatches 0;	Indels 0; Gaps 0;
Oy	1	GATTAGCATTAATAATCTC	20		
Dd	1	GATTAGCATTAATAAATCTC	20		

RESULT 2					
LOCUS	ARI49152/c				
DEFINITION	ARI49152	836 bp	DNA	linear	PAT 08-AUG-2001
ACCESSION	Sequence 6 from patent US 6228579.				
VERSION	ARI49152				
KEYWORDS	ARI49152.1	GI:15113743			
SOURCE	Unknown.				
ORGANISM	Unknown.				
REFERENCE	Unclassified.				
AUTHORS	1 (bases 1 to 836)				
TITLE	Zyskind,J.W. and Forsyth,R.Allyn.				
JOURNAL	Method for identifying microbial proliferation genes				
FEATURES	Patent: US 6228579-A 6 08-Mar-2001;				
source	location/Qualifiers				
	1..836				

RESULT				PAT 08-AUG-2001
3				
Locus	ARI49153			
Definition	Sequence AR149153	836 bp	DNA	
Accession	AR149153			linear
Version	AR149153.1			
Keywords	GI:15113744			
Source	.			
Organism	Unknown:			
Reference	Unclassified.			
Authors	I (bases 1 to 836)			
Title	Zyskind,J.W. and Forsyth,R.Allyn.			
Journal	Method for identifying microbial proliferation genes			
Features	Patent: US 6228579-A 7 Oct-May-2001;			
source	/location/Oqualifiers 1..836			
BASE COUNT	195 a 198 c 220 g 223 t			
ORIGIN	"organism=unknown"			

RESULT 4	
ECOENVA/c	ECOENVA
LOCUS	2048 bp ss-DNA
DEFINITION	linear BCT 20-DEC-1992
ACCESSION	E.coli cell permeability-cell separation protein (enva) gene, complete cds., ftsz gene, 3' end, and secA gene, 5' end. M19211

VERSION	MJ9211.1	GI:145846
KEYWORDS	cell permeability; cell separation protein; envA gene; ftsZ gene; sea gene.	
SOURCE	Escherichia coli (strain K-12) (clone: pACYC184.) DNA.	
ORGANISM	Escherichia coli	
REFERENCE	Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae; Escherichia.	
AUTHORS	1 (bases 1 to 2048)	
TITLE	Beall, B. and Lutkenhaus, J.	
JOURNAL	Sequence analysis, transcriptional organization, and insertional mutagenesis of the envA gene of Escherichia coli	
MEDLINE	J. Bacteriol. 169 (12), 5408-5415 (1987)	
COMMENT	88058745	
FEATURES	Draft entry and computer readable of sequence (1) kindly provided by J. Lutkenhaus (19-FEB-1988). (put. see Kay for more info.).	
SOURCE	Location/Qualifiers	
	1..2048	

Db 1855 GATTACCATATAAATCTC 1836

RESULT 5
LOCUS E27365/c
DEFINITION Treponema pallidum-fused DNA sequence and method for expressing T.
pallidum antigen with the use of the said sequence.
ACCESSION E27365
VERSION E27365.1 GI:13018177
KEYWORDS JP 1999192089-A/4.
SOURCE unidentified.
ORGANISM unidentified.

REFERENCE 1 (bases 1 to 2754)
AUTHORS Katsuya, F., Y.H.H. and Ito.
TITLE Treponema pallidum-fused DNA sequence and method for expressing T.
JOURNAL Patent: JP 1999192089-A 4 21-JUL-1999.
COMMENT FUJIREBIO INC
OS Unidentified
PN JP 1999192089-A/4
PD 21-JUL-1999
PE 29-DEC-1997 JP 1997367638
PR
PI KATSUYA FUJIMURA, YASUHIRO HARA, SATOSHI ITO
PC C12N15/09/C07K14/20/C07K19/00,C12N1/21,C12P21/02,G01N33/53,
PC G01N33/577,
PC (C12N15/09,C12R1:01),(C12N1/21,C12R1:19),(C12P21/02,C12R1:19),
PC C12N15/00,
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CC Strandedness: Double;
CC Topology: Linear;
FH Key 1. 2754 Location/Qualifiers
FT source /organism='unidentified'.
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Source 1. 2754 Location/Qualifiers
1. 2754 /organism='unidentified'
/db_xref='taxon:32644'

BASE COUNT 729 a 680 c 761 g 584 t

ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 2754;
Best Local Similarity 100.0%; Pred. No. 83;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GATTACCATATAAATCTC 20
|||||
Db 49 GATTACCATATAAATCTC 30

RESULT 6
LOCUS ECOSECA
DEFINITION Escherichia coli SecA protein gene, complete cds; mult 5' end.
ACCESSION M20791
VERSION M20791.1 GI:147792
KEYWORDS secA protein.
SOURCE E.coli (strain MC4100) DNA.
ORGANISM Escherichia coli
Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
Escherichia.
1. (bases 1 to 3811)
Schmidt, M., Rolio, E., Grodberg, I. and Oliver, D.
Nucleotide sequence of secA gene and secA(ts) mutations preventing
protein export in Escherichia coli
J. Bacteriol. 170, 3404-3414 (1988)
88298644
JOURNAL
MEDLINE
COMMENT Draft entry and computer readable sequence [1] kindly submitted by
M. Schmidt 28-SEP-1988
The mult gene was identified in Mcl. Gen. Genet. 206, 9-16 (1987)

FEATURES
Source accession number X04831.
Location/Qualifiers
1. 3811
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317..760
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MPAPAKRENAVADITYGTNNIEGFDYLDONMAFSPEERQRLKHALVDEVDSTILDE
NATPLIIGSPADSSSEMYKRVKKIIPHLIRKEDESEFQGGHFSVDEKSNQVNLTE
RGLVLIIEELVKEGIMDEGESLYSPANIMLMHVAALRAHALFTRDVYIVKDEVI
IYDEHTGRMGRMSDGLHQAVERKEGVQIONEQNTASTITFQNFVFLYERLAKMTG
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PVLNGTISTEKSELYSNELTRAGTICAHNLKMFHNAEAIYVAGYAPAAVTATPMAG
RGTDIVLGSGWQAEVALBNPTAEDITEIKADMOYRHDVLEAGCLHITIGBERHSRK
IDMQLRGSRGQDGSSEFYLSMEDALMRIFASRVSGMMKRLGMRGEALIEHPSWT
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VFKATIDAVIPPOSLEEMMDIPGLOERLKNDELDLPIAEMLDKPELHELTRDGL
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BASE COUNT 1016 a 938 c 1004 g 853 t

ORIGIN

Query Match 100.0%; Score 20; DB 1; Length 3811;
Best Local Similarity 100.0%; Pred. No. 78;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GATTACCATATAAATCTC 20
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Db 830 GATTACCATATAAATCTC 811

RESULT 7
LOCUS AE000119/c
DEFINITION Escherichia coli K12 MG1655 section 9 of 400 of the complete
genome.
ACCESSION AE000119 U000096
VERSION AE000119.1 GI:1786283
KEYWORDS
SOURCE Escherichia coli K12.
ORGANISM Escherichia coli K12.
Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
Escherichia.

REFERENCE
AUTHORS
1 (bases 1 to 12434)
Blattner, F.R., Plunkett, G. III, Bloch, C.A., Perna, N.T., Burland, V.,
Bleyer, M., Collado-Vides, J., Glasner, J.D., Rode, C.K., Mayhew, G.F.,
Gregor, J.J., Davis, N.M., Kirkpatrick, H.A., Goeden, M.A., Rose, D.J.,
Mau, B. and Sho, Y.
The complete genome sequence of *Escherichia coli* K-12
Science 277 (5331), 1453-1474 (1997)
97426617
PUBMED
9278503
REFERENCE
AUTHORS
2 (bases 1 to 12434)
Blattner, F.R.
Direct Submission
Submitted (16-JAN-1997) Guy Plunkett III, Laboratory of Genetics,
University of Wisconsin, 445 Henry Mall, Madison, WI 53706, USA.
Email: ecoligenetics.wisc.edu Phone: 608-262-2534 Fax:
608-263-7459
3 (bases 1 to 12434)
Blattner, F.R.
Direct Submission
Submitted (02-SEP-1997) Guy Plunkett III, Laboratory of Genetics,
University of Wisconsin, 445 Henry Mall, Madison, WI 53706, USA.
Email: ecoligenetics.wisc.edu Phone: 608-262-2534 Fax:
608-263-7459
4 (bases 1 to 12434)
Plunkett, G. III.
Direct Submission
Submitted (13-OCT-1998) Laboratory of Genetics, University of
Wisconsin, 445 Henry Mall, Madison, WI 53706, USA
This sequence was determined by the E. coli Genome Project at the
University of Wisconsin-Madison (Frederick R. Blattner, director).
Supported by NIH grants HG00301 and HG01428 (from the Human Genome
Project and NCHGR). The entire sequence was independently
determined from E. coli K12 strain MG1655. Predicted open reading
frames were determined using Genemark software, kindly supplied by
Mark Borodovsky, Georgia Institute of Technology, Atlanta, GA.
30332 (e-mail: markborov@gatech.edu). Open reading frames that
have been correlated with genetic loci are being annotated with CG
Site Nos., unique ID nos. for the genes in the E. coli Genetic
Stock Center (CGSC) database at Yale University, kindly supplied by
Mary Berlyn. A public version of the database is accessible
(http://cgsc.biology.yale.edu). Annotation of the genome is an
ongoing task whose goal is to make the genome sequence more useful
by correlating it with other data. Comments to the authors are
appreciated. Updated information will be available at the E. coli
Genome Project's World Wide Web site
(http://www.genetics.wisc.edu). ** The E. coli K12 sequence and
its annotations are periodically updated; this is version M4. No
sequence changes. Annotation updates: updated gene identifications
and products; all new functional assignments courtesy of Monica
Riley; added promoters, protein binding sites, and repeated
sequences described in reference 1. The unique numeric identifiers
beginning with a lowercase 'b' assigned to each gene (protein- or
RNA-encoding) are now designated as gene synonyms instead of
labels. This should allow them to be searched for in Entrez as gene
names.

FEATURES
source
Location/Qualifiers
1. .12434
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TVGNTIRAFASDNATVYIGTSLDPMDELVTAVAGICDQKREPETLVYNNKOVOOP
VMDRYOOHGMAPLTOEQKPVAKVNVNDNAPQAKRPDYLDIPAPLRKQAD"
1245. .1272
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1313. .2230
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FNHPAIDSSNORVAFSADAFMRQISARFFGMRDIEYLOSGLGSGFDCAIYV
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/function="transport; Protein, peptide secretion"
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/codon_start=1
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gene
promoter
CDS

TITLE	JOURNAL
MEDLINE	PUBMED
REFERENCE	AUTHORS
Genome sequence of enterohaemorrhagic Escherichia coli O157:H7 Nature 409 (6819), 529-533 (2001)	21074935 11206551
(bases 1 to 12518) Perna,N.T., Plunkett,G. III, Burland,V., Mau,B., Glasner,J.D., Rose,D.J., Mayhew,G.F., Evans,P.S., Gregor,J., Kirkpatrick,H.A., Postal,G., Hackett,J., Klink,S., Boutin,A., Shao,Y., Miller,L., Großbeck,E.D., Davis,N.W., Llm,A., Dimantanta,E., Potamouis,K., Apodaca,J., Anantharaman,T.S., Lin,J., Yen,G., Schwartz,D.C., Welch,R.A. and Blattner,F.R.	2 (bases 1 to 12518) Submitted (22-OCT-2000) Laboratory of Genetics, University of Wisconsin, 445 Henry Mall, Madison, WI 53706, USA
Location/Outils	1..12518
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2428..3258	
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2428..3258	
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- REFERENCE 3 (bases 7316 to 10074)
AUTHORS Nakamura, M., Maruyama, I.N., Soma, M., Kato, J.I., Suzuki, H. and Hirota, Y.
TITLE On the process of cellular division in *Escherichia coli*: Nucleotide sequence of the gene for penicillin-binding protein 3
JOURNAL Mol. Gen. Genet. 191, 1-9 (1983)
MEDLINE 83296957
REFERENCE 4 (bases 19464 to 21952)
AUTHORS Robinson, A.C., Kennan, D.J., Hatfull, G.F., Sullivan, N.F., Spiegelberg, R. and Donachie, W.D.
TITLE DNA sequence and transcriptional organization of essential cell division genes *ftsQ* and *ftsA* of *Escherichia coli*: Evidence for overlapping transcriptional units
J. Bacteriol. 160, 546-555 (1984)
MEDLINE 85054557
REFERENCE 5 (bases 21464 to 23333)
AUTHORS Yi, Q.M. and Lutkenhaus, J.
TITLE The nucleotide sequence of the essential cell-division gene *ftsZ* of *Escherichia coli*
Gene 36, 241-247 (1985)
JOURNAL MEDLINE 86083166
REFERENCE 6 (bases 1799 to 2187)
AUTHORS Haughn, G.W., Squitres, C.H., Defelice, M., Largo, C.T. and Calvo, J.M.
TITLE Unusual organization of the *llyA* promoter of *Escherichia coli*
J. Bacteriol. 163, 186-198 (1985)
MEDLINE 85234358
REFERENCE 7 (bases 268 to 1130)
AUTHORS Haughn, G.W., Wessler, S.R., Gemmill, R.M. and Calvo, J.M.
TITLE High A + T content conserved in DNA sequences upstream of *leuABCD* in *Escherichia coli* and *Salmonella typhimurium*
J. Bacteriol. 166, 1113-1117 (1986)
MEDLINE 86223773
REFERENCE 8 (bases 18619 to 19770)
AUTHORS Robinson, A.C., Kennan, D.J., Sweeney, J. and Donachie, W.D.
TITLE Further Evidence for Overlapping Transcriptional Units in an *Escherichia coli* Cell Envelope-Cell Division Gene Cluster: DNA Sequence and Transcriptional Organization of the *ddl ftsQ* Region
J. Bacteriol. 167, 809-817 (1986)
MEDLINE 86304170
REFERENCE 9 (bases 22964 to 25011)
AUTHORS Beall, B. and Lutkenhaus, J.
TITLE Sequence analysis, transcriptional organization, and insertional mutagenesis of the *envA* gene of *Escherichia coli*
J. Bacteriol. 169, 5408-5415 (1987)
MEDLINE 88058745
REFERENCE 10 (bases 27412 to 28277)
AUTHORS Akiyama, M., Horituchi, T. and Sekiguchi, M.
TITLE Molecular cloning and nucleotide sequence of the *murJ* mutator of *Escherichia coli* that causes A:T to C:G transversion
Mol. Gen. Genet. 206, 9-16 (1987)
JOURNAL MEDLINE 87201091
REFERENCE 11 (bases 843 to 1812)
AUTHORS Henikoff, S., Haughn, G.W., Calvo, J.M. and Wallace, J.C.
TITLE A large family of bacterial activator proteins
Proc. Natl. Acad. Sci. U.S.A. 85, 6602-6606 (1988)
MEDLINE 88320486
REFERENCE 12 (bases 20513 to 21772)
AUTHORS Robinson, A.C., Beggs, K.J. and Donachie, W.D.
TITLE Mapping and characterization of mutants of the *Escherichia coli* cell division gene, *ftsA*
Mol. Microbiol. 2, 581-588 (1988)
JOURNAL MEDLINE 89039246
REFERENCE 13 (bases 23989 to 27799)
AUTHORS Schmidt, M., Rollo, E., Grodberg, I. and Oliver, D.
TITLE Nucleotide sequence of *seca* gene and *seca*(ts) mutations preventing protein export in *Escherichia coli*
J. Bacteriol. 170, 3404-3414 (1988)
MEDLINE 88298644
REFERENCE 14 (bases 11142 to 12634)
AUTHORS Parquet, C., Fluore, B., Mengin-Lecreulx, D. and Van Heijenoort, J.
TITLE Nucleotide sequence of the *murF* gene encoding the UDP-MurNac-pentapeptide synthetase of *Escherichia coli*
Nucleic Acids Res. 17, 5379-5379 (1989)
- MEDLINE 89345095
REFERENCE 15 (bases 14743 to 16239)
AUTHORS Ikeda, M., Sato, T., Wachi, M., Jung, H.K., Ishino, F., Kobayashi, M. and Matsunashi, M.
TITLE Structural similarity among *Escherichia coli* *ftsW* and *RodA* proteins and *Bacillus subtilis* SpoVE protein, which function in cell division, cell elongation, and spore formation, respectively
J. Bacteriol. 171, 6375-6378 (1989)
MEDLINE 90036736
REFERENCE 16 (bases 1 to 28277)
AUTHORS Tao, J.S. and Ishiguro, E.E.
TITLE Nucleotide sequence of the *murE* gene of *Escherichia coli*
Can. J. Microbiol. 35, 1051-1054 (1989)
JOURNAL MEDLINE 90124047
REFERENCE 17 (bases 13392 to 15020)
AUTHORS Mengin-Lecreulx, D. and van Heijenoort, J.
TITLE Nucleotide sequence of the *murD* gene encoding the UDP-MurNac-L-Ala-D-Glu synthetase of *Escherichia coli*
Nucleic Acids Res. 18 (1), 183 (1990)
MEDLINE 90174916
REFERENCE 18 (bases 12423 to 15030)
AUTHORS Ikeda, M., Wachi, M., Ishino, F. and Matsunashi, M.
TITLE Nucleotide sequence involving *murD* and an open reading frame *orf-Y* spacing *murF* and *ftsW* in *Escherichia coli*
Nucleic Acids Res. 18, 1058-1058 (1990)
JOURNAL MEDLINE 90192099
REFERENCE 19 (bases 6088 to 7587)
AUTHORS Gomez, M.J., Fluore, B., Van Heijenoort, J. and Ayala, J.A.
TITLE Nucleotide sequence of the regulatory region of *pbbp* gene of *Escherichia coli*
Nucleic Acids Res. 18, 2813-2813 (1990)
MEDLINE 90251464
REFERENCE 20 (bases 4274 to 6093)
AUTHORS Leclerc, G., Noel, G. and Drapeau, G.
TITLE Molecular cloning, nucleotide sequence and expression of *shl*, a new gene in the 2-minute region of the genetic map of *Escherichia coli*
J. Bacteriol. 172, 4696-4700 (1990)
MEDLINE 90330585
REFERENCE 21 (bases 16094 to 18886)
AUTHORS Ikeda, M., Wachi, M., Jung, H.K., Ishino, F. and Matsunashi, M.
TITLE Nucleotide sequence involving *murG* and *murC* in the *mra* gene cluster region of *Escherichia coli*
Nucleic Acids Res. 18, 4014-4014 (1990)
JOURNAL MEDLINE 90326550
REFERENCE 22 (bases 16094 to 17806)
AUTHORS Mengin-Lecreulx, D., Texier, L. and van Heijenoort, J.
TITLE Nucleotide sequence of the cell-envelope *murG* gene of *Escherichia coli*
Nucleic Acids Res. 18 (9), 2810 (1990)
JOURNAL MEDLINE 90251461
REFERENCE 23 (bases 1 to 28277)
AUTHORS Michaud, C., Parquet, C., Fluore, B., Blanot, D. and van Heijenoort, J.
TITLE Revised interpretation of the sequence containing the *murE* gene encoding the UDP-N-acetylmuramyl-tripeptide synthetase of *Escherichia coli*
Biochem. J. 269 (1), 277-278 (1990)
MEDLINE 90328986
REFERENCE 24 (bases 1 to 28277)
AUTHORS Wang, Q. and Calvo, J.M.
TITLE Lrp, a global regulatory protein of *Escherichia coli* binds co-operatively to multiple sites and activates transcription of *llyA*
J. Mol. Biol. 229, 306-318 (1993)
JOURNAL MEDLINE 93156044
REFERENCE 25 (bases 1 to 28277)
AUTHORS Wang, Q. and Calvo, J.M.
TITLE Lrp, a major regulatory protein in *Escherichia coli* bends DNA and can organize the assembly of a higher-order nucleoprotein structure
EMBO J. 12, 2495-2501 (1993)
MEDLINE 93285120
REFERENCE 26 (bases 4274 to 6093)
AUTHORS Jahreis, K., Postma, P.W. and Lengeler, J.W.
TITLE Nucleotide sequence of the *lly H-fur* gene of *Escherichia coli* K-12

JOURNAL	unpublished	and Salmonella typhimurium LT2
REFERENCE	27 (bases 1 to 28277)	
AUTHORS	Ayala,J.A.	
TITLE	Regulation of transcription at the 2-minute region of the genetic map of Escherichia coli	
JOURNAL	unpublished	
REFERENCE	28 (bases 1 to 28277)	
AUTHORS	Ayala,J.A.	
TITLE	Direct Submission	
JOURNAL	Submitted (08-JAN-1991)	
REFERENCE	Molecular, Centro de Biologia Molecular, Universidad Autonoma, Canto-Blanco 28049, Madrid, Spain	
COMMENT	This entry comprises a merged sequence of 28kb of which a portion is the submittor's original work.	
FEATURES	Location/Qualifiers	
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		
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Db 24817	GATTAGCATATAAATCTC 24798	
RESULT 10		
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DEFINITION	AX191720	28277 bp DNA linear PAT 15-AUG-2001
ACCESSION	AX191720	
VERSION	AX191720.1	GI:15209889
KEYWORDS		
SOURCE		
ORGANISM	Escherichia coli.	
	Escherichia coli	
	Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae; Escherichia.	
REFERENCE	1 (bases 1 to 28277)	
AUTHORS	Iversen,P.L.	
TITLE	Antisense antibacterial cell division composition and method	
JOURNAL	Patent: WO 0149775-A 2 12-JUL-2001;	
	Avi Biopharma, Inc. (US)	
FEATURES	Location/Qualifiers	
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Best Local Similarity	100.0%; Pred. NO. 52;	
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		
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Db 24817	GATTAGCATATAAATCTC 24798	
RESULT 11		

LOCUS	ECOL10K	111401 bp	DNA	linear	BCT 27-JUN-1997
DEFINITION	E.coli K12 genome, 0-2.4mu, region.				
ACCESSION	D10483 J01597 J01683 J01706 K01288 M10420 M10611 M12544				
VERSION	V00259 X04711 X54847 X54945 X55034 X56742				
KEYWORDS	D10483.1 GI:216434				
SOURCE	Gen. Dna.				
ORGANISM	Escherichia coli (strain:K-12) DNA.				
REFERENCE	Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae; Escherichia.				
AUTHORS	1 (bases 1 to 111401)				
TITLE	Mori,H.				
JOURNAL	Direct Submission				
REFERENCE	Submitted (18-FEB-1992) Hirotsada Mori, Institute for Virus Research, Kyoto University, Genetics and Molecular Biology, 53 Shogoin Kawara-Machi, Sakyo-ku, Kyoto 606, Japan				
AUTHORS	(E-mail:es2985@sakura.kudpc.kyoto-u.ac.jp, Tel:075-751-4042, Fax:075-761-5626)				
TITLE	2 (sites)				
AUTHORS	Smith,B.R. and Schleif,R.				
REFERENCE	Nucleotide sequence of the L-arabinose regulatory region of Escherichia coli K12				
JOURNAL	The Journal of Biological Chemistry. 253 (19), 6931-6933 (1978)				
DEFINITION	79005683				
ACCESSION	357433				
VERSION	3 (sites)				
KEYWORDS	Ohtsubo,H. and Ohtsubo,E.				
SOURCE	Nucleotide sequence of an insertion element, IS1				
ORGANISM	Proceedings of the National Academy of Sciences of the United States of America. 75 (2), 615-619 (1978)				
REFERENCE	78137003				
AUTHORS	273224				
TITLE	4 (sites)				
JOURNAL	Greenfield,L., Boone,T. and Wilcox,G.				
DEFINITION	DNA sequence of the araBAD promoter in Escherichia coli B/r				
ACCESSION	Proceedings of the National Academy of Sciences of the United States of America. 75 (10), 4724-4728 (1978)				
VERSION	79116194				
KEYWORDS	368797				
SOURCE	5 (sites)				
ORGANISM	Johnsrud,L.				
REFERENCE	DNA sequence of the transposable element IS1				
JOURNAL	Molecular & general genetics : MGG. 169 (2), 213-218 (1979)				
DEFINITION	79177885				
ACCESSION	375010				
VERSION	6 (sites)				
KEYWORDS	Smith,D.R. and Calvo,J.M.				
SOURCE	Nucleotide sequence of the E coli gene coding for dihydrofolate reductase				
ORGANISM	Nucleic acids research. 8 (10), 2255-2274 (1980)				
REFERENCE	81053692				
AUTHORS	6159575				
TITLE	7 (sites)				
JOURNAL	Myadec,G.G., Horwitz,A.H., Cass,L.G., Timko,J. and Wilcox,G.				
DEFINITION	DNA sequence of the araC regulatory gene from Escherichia coli B/r				
ACCESSION	Nucleic acids research. 8 (22), 5267-5274 (1980)				
VERSION	81124262				
KEYWORDS	7008027				
SOURCE	8 (sites)				
ORGANISM	Ogden,S., Hagerly,D., Stoner,C.M., Kolodrubetz,D. and Schleif,R.				
REFERENCE	The Escherichia coli L-arabinose operon: binding sites of the regulatory proteins and a mechanism of positive and negative regulation				
JOURNAL	Proceedings of the National Academy of Sciences of the United States of America. 77 (6), 3346-3350 (1980)				
DEFINITION	81013881				
ACCESSION	6251457				
VERSION	9 (sites)				
KEYWORDS	Katinka,M., Cossart,P., Sibillii,L., Saint-Gironis,I., Chailivnac,M.A., Le Bris,G., Cohen,G.N. and Taitiv,M.				
SOURCE	Nucleotide sequence of the thrA gene of Escherichia coli				
ORGANISM	Proceedings of the National Academy of Sciences of the United States of America. 77 (6), 3346-3350 (1980)				

MEDLINE 81077247
PUBMED 7003595
REFERENCE 10 (sites)
AUTHORS Mackie,G.A.
TITLE Nucleotide sequence of the gene for ribosomal protein S20 and its flanking regions
JOURNAL The Journal of biological chemistry. 256 (15), 8177-8182 (1981)
MEDLINE 81264207
PUBMED 6267039
REFERENCE 11 (sites)
AUTHORS Cossart,P., Kalinka,M. and Yaniv,M.
TITLE Nucleotide sequence of the thrB gene of E. coli, and its two adjacent regions: the thrAB and thrBC junctions
JOURNAL Nucleic acids research. 9 (2), 339-347 (1981)
MEDLINE 81150470
PUBMED 6259626
REFERENCE 12 (sites)
AUTHORS Lee,N.L., Gielow,W.O. and Wallace,R.G.
TITLE Mechanism of araC autoregulation and the domains of two overlapping promoters, Pc and PBAD, in the L-arabinose regulatory region of Escherichia coli
JOURNAL Proceedings of the National Academy of Sciences of the United States of America. 78 (2), 752-756 (1981)
MEDLINE 81199399
PUBMED 6262769
REFERENCE 13 (sites)
AUTHORS Stoner,C.M. and Schleif,R.
TITLE Is the amino acid but not the nucleotide sequence of the Escherichia coli araC gene conserved?
JOURNAL Journal of molecular biology. 154 (4), 649-652 (1982)
MEDLINE 82216830
PUBMED 6283093
REFERENCE 14 (sites)
AUTHORS Gilson,E., Nikaido,H. and Hofnung,M.
TITLE Sequence of the malK gene in E.coli K12
JOURNAL Nucleic acids research. 10 (22), 7449-7458 (1982)
MEDLINE 83116968
PUBMED 6296778
REFERENCE 15 (sites)
AUTHORS Parsot,C., Cossart,P., Saint-Girons,I. and Cohen,G.N.
TITLE Nucleotide sequence of thrC and of the transcription termination region of the threonine operon in Escherichia coli K12
JOURNAL Nucleic acids research. 11 (21), 7331-7345 (1983)
MEDLINE 84089770
PUBMED 6316258
REFERENCE 16 (sites)
AUTHORS Bouvier,J., Richard,C., Richard,F., Patte,J.C. and Stragier,P.
TITLE Nucleotide sequence and expression of the Escherichia coli dcpB gene
JOURNAL The Journal of biological chemistry. 259 (23), 14829-14834 (1984)
MEDLINE 85054974
PUBMED 6094578
REFERENCE 17 (sites)
AUTHORS Bardwell,J.C. and Craig,E.A.
TITLE Major heat shock gene of Drosophila and the Escherichia coli heat-inducible dnaK gene are homologous
JOURNAL Proceedings of the National Academy of Sciences of the United States of America. 81 (3), 848-852 (1984)
MEDLINE 84144800
PUBMED 6322174
REFERENCE 18 (sites)
AUTHORS Innis,M.A., Tokunaga,M., Williams,M.E., Lorange,J.M., Chang,S.Y., Chang,S. and Wu,H.C.
TITLE Nucleotide sequence of the Escherichia coli prolipoprotein signal peptidase (lsp) gene
JOURNAL Proceedings of the National Academy of Sciences of the United States of America. 81 (12), 3708-3712 (1984)
MEDLINE 84222028
PUBMED 6374664
REFERENCE 19 (sites)
AUTHORS Bouvier,J., Patte,J.C. and Stragier,P.
TITLE Multiple regulatory signals in the control region of the

JOURNAL Escherichia coli carAB operon
PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF AMERICA. 81 (13), 4139-4143 (1984)
MEDLINE 84248073
PUBMED 6377309
REFERENCE 20 (sites)
AUTHORS Chong,P., Hui,I., Loo,T. and Gillam,S.
TITLE Structural analysis of a new GC-specific insertion element IS186
JOURNAL FEBS letters. 192 (1), 47-52 (1985)
MEDLINE 86030702
PUBMED 2996940
REFERENCE 21 (sites)
AUTHORS Kamio,Y., Lin,C.K., Regue,M. and Wu,H.C.
TITLE Characterization of the lles-lsp operon in Escherichia coli. Identification of an open reading frame upstream of the lles gene and potential promoter(s) for the lles-lsp operon
JOURNAL The Journal of biological chemistry. 260 (9), 5616-5620 (1985)
MEDLINE 85182715
PUBMED 2985604
REFERENCE 22 (sites)
AUTHORS Friedberg,D., Rosenthal,E.R., Jones,J.W. and Calvo,J.M.
TITLE Characterization of the 3' end of the leucine operon of Salmonella typhimurium
JOURNAL Molecular & general genetics : MGG. 199 (3), 486-494 (1985)
MEDLINE 85295470
PUBMED 2993799
REFERENCE 23 (sites)
AUTHORS Cowling,D.W., Bardwell,J.C., Craig,E.A., Woolford,C., Hendrix,R.W. and Gross,C.A.
TITLE Consensus sequence for Escherichia coli heat shock gene promoters
JOURNAL Proceedings of the National Academy of Sciences of the United States of America. 82 (9), 2679-2683 (1985)
MEDLINE 85190560
PUBMED 3887408
REFERENCE 24 (sites)
AUTHORS Sekiguchi,T., Ortega-Cesena,J., Nosoh,Y., Ohashi,S., Tsuda,K. and Kanaya,S.
TITLE DNA and amino-acid sequences of 3-isopropylmalate dehydrogenase of Bacillus coagulans. Comparison with the enzymes of Saccharomyces cerevisiae and Thermus thermophilus
JOURNAL Biochim. Biophys. Acta. 867, 36-44 (1986)
MEDLINE 87163495
PUBMED 3549454
REFERENCE 25 (sites)
AUTHORS Lee,N., Gielow,W., Martin,R., Hamilton,E. and Fowler,A.
TITLE The organization of the arabid operon of Escherichia coli
JOURNAL Gene. 47 (2-3), 231-244 (1986)
MEDLINE 86111849
PUBMED 3003084
REFERENCE 26 (sites)
AUTHORS Ohki,M., Tamura,F., Nishimura,S. and Uchida,H.
TITLE Nucleotide sequence of the Escherichia coli dnaJ gene and purification of the gene product
JOURNAL The Journal of biological chemistry. 261 (4), 1778-1781 (1986)
MEDLINE 86111849
PUBMED 3003084
REFERENCE 27 (sites)
AUTHORS

Query Match 100.0%; Score 20; DB 1; Length 111401;
Best Local Similarity 100.0%; Pred. No. 40;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTAGCATTAATAATCTC 20
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Db 107941 GATTAGCATTAATAATCTC 107922

RESULT 12
AC055113
LOCUS 155976 bp DNA linear HTG 24-ANG-2000
DEFINITION Homo sapiens chromosome 12 clone CTD-2021H9, WORKING DRAFT
AC055113
AC055113.2 GI:9838029
HTG: HTGS_PHASE1; HTGS_DRAFT.

SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 155976)
AUTHORS Waterston,R.H.
TITLE The sequence of Homo sapiens clone
JOURNAL Unpublished
2 (bases 1 to 155976)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (17-APR-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Aug 17, 2000 this sequence version replaced gi:7579846.
COMMENT ----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: MUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H_MS2021H09
----- Summary Statistics -----
Sequencing vector: M13; 100%
Sequencing vector: plasmid; 0%
Chemistry: Dye-primer ET; 10% of reads
Chemistry: Dye-terminator Big Dye; 0% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 14393 bases at least Q40
Consensus quality: 146482 bases at least Q30
Consensus quality: 147905 bases at least Q20
Insert size: 18000; agarose-fp
Insert size: 154176; sum-of-ctlgis
Quality coverage: 5.12 in Q20 bases; agarose-fp
Quality coverage: 5.01 in Q20 bases; sum-of-ctlgis

* NOTE: This is a 'working draft' sequence. It currently
* consists of 19 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 3520: contig of 3520 bp in length
* 3521 3620: gap of unknown length
* 3621 9066: contig of 5446 bp in length
* 9067 9166: gap of unknown length
* 9167 13195: contig of 4029 bp in length
* 13196 13295: gap of unknown length
* 13296 17854: contig of 4559 bp in length
* 17855 17954: gap of unknown length
* 17955 26904: contig of 8950 bp in length
* 26905 27004: gap of unknown length
* 27005 35606: contig of 8602 bp in length
* 35607 35706: gap of unknown length
* 35707 43003: contig of 7297 bp in length
* 43004 43103: gap of unknown length
* 43104 52582: contig of 9479 bp in length
* 52583 52582: gap of unknown length
* 52583 62116: contig of 9434 bp in length
* 62117 62216: gap of unknown length
* 62217 75369: contig of 13153 bp in length
* 75370 75469: gap of unknown length
* 75470 88516: contig of 13047 bp in length
* 88517 88616: gap of unknown length
* 88617 104061: contig of 15445 bp in length
* 104062 104161: gap of unknown length
* 104162 119796: contig of 15635 bp in length
* 119797 119896: gap of unknown length
* 119897 141280: contig of 21384 bp in length
* 141281 141380: gap of unknown length
* 141381 142833: contig of 1453 bp in length

FEATURES
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/chromosome="12"
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3621. 9066
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9167. 13195
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13296. 17854
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17955. 26904
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27005. 35606
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35707. 43003
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43104. 52582
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52683. 62116
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62217. 75369
/note="assembly_name:Contig19"
75470. 88516
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88617. 104061
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104162. 119796
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119897. 141280
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148667. 152318
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152419. 155976
/note="assembly_name:Contig28"
155976. 155976
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1802 others
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Best Local Similarity 100.0%; Pred. No. 98;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 ATTACGATTAATAATCTC 20
Db 94819 ATTACGATTAATAATCTC 94837
RESULT 13
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LOCUS Homo sapiens 12q BAC RP11-158011 (Roswell Park Cancer Institute
DEFINITION Human BAC library) complete sequence.
ACCESSION AC084877
VERSION AC084877.18 GI:14277186
KEYWORDS HTG.
SOURCE human.


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886..1010
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repeat_region
1299..1616
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repeat_region
2804..2898
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repeat_region
3417..3449
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3827..3900
/rpt_family="L2"
repeat_region
4383..4419
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6920..6970
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Best Local Similarity 100.0%; Pred. No. 96;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 10996 ATTAGCATATAAATCTC 10978

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RESULT 14
LOCUS AE008700/c 22286 bp DNA linear BCT 25-OCT-2001
DEFINITION *Salmonella typhimurium* LT2, section 8 of 224 of the complete genome.
ACCESSION AE008700 AE006468
VERSION AE008700.1 GI:16418628
KEYWORDS
SOURCE *Salmonella typhimurium* LT2.
ORGANISM *Salmonella typhimurium* LT2.
Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae; *Salmonella*.
1 (bases 1 to 22286)
McCelland, M., Sanderson, K.E., Spieth, J., Clifton, S.W., Latreille, P., Courtney, U., Portolillo, S., Ali, J., Dante, M., Du, F., Hou, S., Layman, D., Leonard, S., Nguyen, C., Scott, K., Holmes, A.,

JOURNAL
PUBMED
REFERENCE
AUTHORS
TITLE
JOURNAL
2 (bases 1 to 22286)
The *Salmonella typhimurium* Genome Sequencing Project.
Direct Submission
Submitted (29-MAR-2001) Genome Sequencing Center, Department of Genetics, Washington University School of Medicine, 4444 Forest Park Boulevard, St. Louis MO 63108, USA
Supported by NIH grant 5U 01 AI43283

COMMENT

Coding sequences below are predicted from manually evaluated computer analysis, using similarity information and the programs; GLIMMER: <http://www.tigr.org/softlab/glimmer/glimmer.html> and Genemark; <http://opal.biology.gatech.edu/Genemark/>

EC numbers were kindly provided by Junko Yabuzaki and the Kyoto Encyclopedia of Genes and Genomes; <http://www.genome.ad.jp/kegg/>, and Pedro Romero and Peter Karp at EcoCyc; <http://ecocyc.org/PangeaSystems.com/ecocyc/>

The analyses of ribosome binding sites and promoter binding sites were kindly provided by Heladia Salgado, Julio Collado-Vides and ReguionDB; http://kinich.cifn.unam.mx:8850/db/reguiondb_intro.frameset

FEATURES

source
1. 22286
/organism="Salmonella typhimurium LT2"
/strain="LT2; SGSC 1412; ATCC 700720"
/db_xref="ATCC:700720"
/db_xref="taxon:99287"
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84..89
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96..1571
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UDP-N-acetyl-muramate:alanine ligase"
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RBS
gene

CDS /note="STM0130"
1564..2484
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LMGAGLVAAPVATLRAEFEGKLEKQKARISALGLPIVPSREGSVGKTYEE
NALGALSLAFQHDEILIEKWLCEPFTVAIVGEIIPISIRIPAGTEFYDEAYLS
DETQYFCPAGLDAQSEALQSLVLAQMKALGCTGCRIDVMDSDGQFYLLAEANTSPG
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2328..3316
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2328..3316
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2354..3362
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2467..2472
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2486..3316
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YGPESASEVLOGYREMGQVLAKEFTLKEAMPTARSMQTLNNGIKLNLGRGDTMK
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3313..4575
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4636..5787
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ITIPDKLKLVLGRISLIDFANGANDVAGVQVAILITRPGMLNVDVDFVRS
EMGVAMGSSVAGSEDAEAKMAISSPLELIDLSARGVLVNTGFPDRIDEF
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VPSVGRPPRSIQRTLAIEVLEPRTLELNLVNEITLQDLQKQGVKHHLAGVIL
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Best local similarity 95.0%; Pred. No. 2.6e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GATTACGATTAATAATCTC 20
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Db 7650 GATTACGATTAATAAGTCTC 7631

RESULT 15
AC036102/C 127603 bp DNA linear PRI 16-OCT-2001
LOCUS Homo sapiens chromosome 15 clone CTD-2297L20 map 15q21.1, complete
DEFINITION sequence.
AC036102.8 GI:16152267
VERSION AC036102.8
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 127603)
Rosen, L., Madan, A., Qin, S., Baradaran, L., Birditt, B., Bloom, S.,
Burke, J., Dors, M., Fleetwood, P., Kaur, A., Madan, A., Nesbitt, R.,
Pate, D., and Hood, L.
Pate, D., and Hood, L.
Unpublished
2 (bases 1 to 127603)
Rosen, L., Madan, A., Qin, S., Abbasi, N., Baradaran, L., Birditt, B.,
Bloom, S., Dors, M., Dickhoff, R., Fleetwood, P., Harrison, G.,
James, R., Kaur, A., Madan, A., Owen, M. P., Ratcliffe, A., Shaffer, T.,
and Hood, L.

TITLE Direct Submission
JOURNAL Submitted (07-APR-2000) Multimegabase Sequencing Center, University of Washington, PO BOX 357730, Seattle, WA 98195, USA
REFERENCE 3 (bases 1 to 127603)
AUTHORS Rowen, L., Madan, A., Qin, S., Baradarani, L., Birditt, B., Bloom, S., Burke, J., Dots, M., Fleetwood, P., Kaur, A., Madan, A., Nesbitt, R., Pale, D., and Hood, L.
TITLE Direct Submission
JOURNAL Submitted (16-OCT-2001) Multimegabase Sequencing Center, Institute for Systems Biology, 4225 Roosevelt Way NE, Suite 200, Seattle, WA 98105, USA
COMMENT On Oct 16, 2001 this sequence version replaced gi:14318379.
 ----- Genome Center
 Center: Multimegabase Sequencing Center
 Center code: UWMSC
 Web site: http://chroma.mbt.washington.edu/msg_www
 Contact: leetowensystemsbiology.org
 ----- Summary Statistics
 Sequencing vector: pUC18; L08752
 Chemistry: Dye-terminator Big Dye; 90% of reads
 Chemistry: Dye-primer Big Dye; 10% of reads
 Assembly program: Phrap; version 0.990399
 Note: data from AC013452 [Drafting center UWMSC] and AC022306 [Drafting center UWMSC] were added for finishing.
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 /note="This clone overlaps RP11-325E5 AC013452 and RP11-295H24 AC022306. Data from overlapping BACs combined and the consensus sequence determined from CTD-2297L20 to the extent possible."
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 85642
 /note="low quality data."
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 /note="Overlap with RP11-295H24 AC022306."
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BASE COUNT
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 Best Local Similarity 95.0%; Pred. No. 1.8e+02;
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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 DB 61460 GAGTAGCATATTAATCTC 61441
 Search completed: October 6, 2002, 16:01:31
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